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#25	Search tpa AND upa Limits: Publication Date to 1998	16:32:26	315
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#15	Search upa AND leukemia Limits: Publication Date to 1998	16:25:55	26
#8	Related Articles for PubMed (Select 7927776)	15:40:50	134

#11 Search lethal factor and cancer and plasminogen Limits: Publication Date to 1998	14:21:48	2
#10 Search lethal factor and cancer Limits: Publication Date to 1998	14:21:31	377
#9 Search leppla and lethal factor and cancer Field: All Fields , Limits: Publication Date to 1998	14:20:14	1
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Nov 16 2004 07:00:47

FILE 'MEDLINE' ENTERED AT 10:57:42 ON 18 NOV 2004

E LEPPLA S/AU
L1 116 S LEPPLA S?/AU
L2 433 S LETHAL FACTOR
L3 1325 S PROTECTIVE ANTIGEN
L4 188 S L3 AND L2
L5 188 S L4 AND L2
L6 1671881 S CANCER? OR TUMOR? OR NEOPLAS? OR METASTA?
L7 13 S L5 AND L6
L8 27177 S PLASMINOGEN ACTIVATOR
L9 3 S L8 AND L7
L10 188 S L2 AND L3
L11 13 S L10 AND L6
L12 3 S L8 AND L11

FILE 'CANCERLIT' ENTERED AT 11:04:32 ON 18 NOV 2004

9 S LEPPLA S?/AU
L13 22 S LETHAL FACTOR
L14 68 S PROTECTIVE ANTIGEN
L15 5939 S PLASMINOGEN ACTIVATOR
L16 54 S ANTHRAX
L17 11 S L17 AND L14
L18 6646 S PLASMINOGEN
L19 1 S L19 AND L18
L20

FILE 'CAPLUS' ENTERED AT 11:06:04 ON 18 NOV 2004

151 S LEPPLA S?/AU
L21 567 S LETHAL FACTOR
L22 1402 S PROTECTIVE ANTIGEN
L23 25343 S PLASMINOGEN
L24 687350 S CANCER? OR TUMOR? OR NEOPLAS? OR METASTA?
L25 6 S L22 AND L24
L26 5 S L26 AND L25
L27

FILE 'PCTFULL' ENTERED AT 11:07:26 ON 18 NOV 2004

6 S LEPPLA S?/AU
L28 246 S LETHAL FACTOR
L29 8577 S PLASMINOGEN
L30 82388 S CANCER? OR TUMOR? OR NEOPLAS? OR METASTA?
L31 25 S L29 AND L30
L32 24 S L32 AND L31
L33 2 S L33 NOT PY>1999
L34

FILE 'MEDLINE, CANCERLIT, CAPLUS, PCTFULL' ENTERED AT 11:09:56 ON 18 NOV 2004

L35 7 DUP REM L12 L20 L27 L34 (4 DUPLICATES REMOVED)

L35 ANSWER 7 OF 7 PCTFULL COPYRIGHT 2004 Univentio on STN
 ACCESSION NUMBER: 1998049311 PCTFULL ED 20020514
 TITLE (ENGLISH): RICIN-LIKE TOXIN VARIANTS FOR TREATMENT OF
CANCER, VIRAL OR PARASITIC INFECTIONS
 TITLE (FRENCH): VARIANTES DE TOXINES DE TYPE RICIN DESTINEES AU
 TRAITEMENT D'INFECTIONS **CANCEREUSES**, VIRALES
 OU PARASITAIRES
 INVENTOR(S): BORGFORD, Thor
 PATENT ASSIGNEE(S): DE NOVO ENZYME CORPORATION;
 BORGFORD, Thor
 LANGUAGE OF PUBL.: English
 DOCUMENT TYPE: Patent
 PATENT INFORMATION:

NUMBER	KIND	DATE
WO 9849311	A2	19981105

DESIGNATED STATES
 W:

AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE
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 BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF
 BJ CF CG CI CM GA GN ML MR NE SN TD TG

APPLICATION INFO.: WO 1998-CA394 A 19980430
 PRIORITY INFO.: US 1997-60/045,148 19970430
 US 1997-60/063,715 19971029

ANSWER 2 OF 3 MEDLINE on STN
ACCESSION NUMBER: 2003031708 MEDLINE
DOCUMENT NUMBER: PubMed ID: 12525700
TITLE: Potent antitumor activity of a urokinase-activated
engineered anthrax toxin.
AUTHOR: Liu Shihui; Aaronson Hannah; Mitola David J; Leppla Stephen
H; Bugge Thomas H
CORPORATE SOURCE: Oral Infection and Immunity Branch, National Institute of
Dental and Craniofacial Research, National Institutes of
Health, Bethesda, MD 20892, USA.
SOURCE: Proceedings of the National Academy of Sciences of the
United States of America, (2003 Jan 21) 100 (2) 657-62.
Journal code: 7505876. ISSN: 0027-8424.
PUB. COUNTRY: United States
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
LANGUAGE: English
FILE SEGMENT: Priority Journals
ENTRY MONTH: 200302
ENTRY DATE: Entered STN: 20030123
Last Updated on STN: 20030225
Entered Medline: 20030224

ED Entered STN: 20030123

Last Updated on STN: 20030225

Entered Medline: 20030224

AB The acquisition of cell-surface urokinase **plasminogen
activator** activity is a hallmark of malignancy. We generated an
engineered anthrax toxin that is activated by cell-surface urokinase in
vivo and displays limited toxicity to normal tissue but broad and potent
tumoricidal activity. Native anthrax toxin **protective
antigen**, when administered with a chimeric anthrax toxin
lethal factor, *Pseudomonas* exotoxin fusion protein, was
extremely toxic to mice, causing rapid and fatal organ damage. Replacing
the furin activation sequence in anthrax toxin **protective
antigen** with an artificial peptide sequence efficiently activated
by urokinase greatly attenuated toxicity to mice. In addition, the
mutation conferred cell-surface urokinase-dependent toxin activation in
vivo, as determined by using a panel of plasminogen, **plasminogen
activator**, **plasminogen activator** receptor, and
plasminogen activator inhibitor-deficient mice.
Surprisingly, toxin activation critically depended on both urokinase
plasminogen activator receptor and plasminogen in vivo,
showing that both proteins are essential cofactors for the generation of
cell-surface urokinase. The engineered toxin displayed potent
tumor cell cytotoxicity to a spectrum of transplanted
tumors of diverse origin and could eradicate established solid
tumors. This **tumoricidal** activity depended strictly on
tumor cell-surface plasminogen activation. The data show that a
simple change of protease activation specificity converts anthrax toxin
from a highly lethal to a potent **tumoricidal** agent.

L9 ANSWER 3 OF 3 MEDLINE on STN
ACCESSION NUMBER: 2001276184 MEDLINE
DOCUMENT NUMBER: PubMed ID: 11278833
TITLE: Targeting of **tumor** cells by cell surface
urokinase **plasminogen activator**
-dependent anthrax toxin.
AUTHOR: Liu S; Bugge T H; Leppla S H
CORPORATE SOURCE: Oral Infection and Immunity Branch and Oral and Pharyngeal
Cancer Branch, NIDCR, National Institutes of Health,
Bethesda, Maryland 20892, USA.
SOURCE: Journal of biological chemistry, (2001 May 25) 276 (21)
17976-84.
Journal code: 2985121R. ISSN: 0021-9258.
PUB. COUNTRY: United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
LANGUAGE: English
FILE SEGMENT: Priority Journals
ENTRY MONTH: 200107
ENTRY DATE: Entered STN: 20010709
Last Updated on STN: 20030105
Entered Medline: 20010705

ED Entered STN: 20010709

Last Updated on STN: 20030105

Entered Medline: 20010705

AB Urokinase **plasminogen activator** receptor (uPAR) binds pro-urokinase **plasminogen activator** (pro-uPA) and thereby localizes it near plasminogen, causing the generation of active uPA and plasmin on the cell surface. uPAR and uPA are overexpressed in a variety of human **tumors** and **tumor** cell lines, and expression of uPAR and uPA is highly correlated to **tumor** invasion and **metastasis**. To exploit these characteristics in the design of **tumor** cell-selective cytotoxins, we constructed mutated anthrax toxin-**protective antigen** (PrAg) proteins in which the furin cleavage site is replaced by sequences cleaved specifically by uPA. These uPA-targeted PrAg proteins were activated selectively on the surface of uPAR-expressing **tumor** cells in the presence of pro-uPA and plasminogen. The activated PrAg proteins caused internalization of a recombinant cytotoxin, FP59, consisting of anthrax toxin **lethal factor** residues 1-254 fused to the ADP-ribosylation domain of Pseudomonas exotoxin A, thereby killing the uPAR-expressing **tumor** cells. The activation and cytotoxicity of these uPA-targeted PrAg proteins were strictly dependent on the integrity of the **tumor** cell surface-associated plasminogen activation system. We also constructed a mutated PrAg protein that selectively killed tissue **plasminogen activator**-expressing cells. These mutated PrAg proteins may be useful as new therapeutic agents for **cancer** treatment.